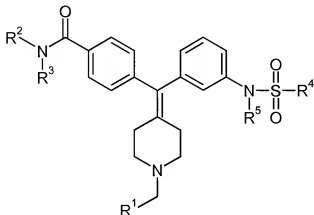


Listing of Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (currently amended) A compound of formula I, or a pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof:



I

wherein

R¹ is phenyl; pyridyl; thienyl; furyl; imidazolyl; pyrrolyl; or thiazolyl, wherein R¹ is optionally substituted with one or more groups selected from C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy, chloro, fluoro, bromo, and iodo; selected from C₆₋₁₀aryl and C₂₋₆heteroaryl, wherein said C₆₋₁₀aryl and C₂₋₆heteroaryl are optionally substituted with one or more groups selected from -R, -NO₂, -OR, -Cl, -Br, -I, -F, -CF₃, -C(=O)R, -C(=O)OH, -NH₂, -SH, -NHR, -NR₂, -SR, -SO₃H, -SO₂R, -S(=O)R, -CN, -OH, -C(=O)OR, -C(=O)NR₂, -NRC(=O)R, and -NRC(=O)-OR, wherein R is, independently, a hydrogen or C₁₋₆alkyl; and R², R³, and R⁴ and R⁵ are, independently, C₁₋₃alkyl or halogenated C₁₋₃alkyl selected from hydrogen, C₁₋₆alkyl, and C₃₋₆cycloalkyl, wherein said C₁₋₆alkyl and C₃₋₆cycloalkyl are optionally substituted with one or more groups selected from -R, -NO₂, -OR, -Cl, -Br, -I, -F, -CF₃, -C(=O)R, -C(=O)OH, -NH₂, -SH, -NHR, -NR₂, -SR, -SO₃H, -SO₂R, -S(=O)R, -CN, -OH, -C(=O)OR, -C(=O)NR₂, -NRC(=O)R, and -NRC(=O)-OR, wherein R is, independently, a hydrogen or C₁₋₆alkyl; and R⁵ is hydrogen.

2. (currently amended) A compound according to claim 1, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof

wherein R¹ is ~~selected from~~ phenyl; pyridyl; thienyl; furyl; ~~imidazolyl; triazolyl; pyrrolyl; or~~ thiazolyl; and N-oxide-pyridyl, wherein R⁴ is ~~optionally substituted with one or more groups selected from~~ C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy, chloro, fluoro, bromo, and iodo;

R², R³, and R⁴ are, independently, C₁₋₃alkyl or halogenated C₁₋₃alkyl; and

R⁵ is selected from hydrogen, C₁₋₆alkyl, and ~~or~~ C₃₋₆cycloalkyl, wherein said C₁₋₆alkyl and C₃₋₆cycloalkyl are ~~optionally substituted with one or more groups selected from~~ C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy, chloro, fluoro, bromo, and iodo.

3. (currently amended) A compound according to claim 2 ~~claim 4, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof~~

wherein R¹ is ~~selected from~~ phenyl; pyridyl; thienyl; furyl; ~~imidazolyl; pyrrolyl; and or~~ thiazolyl, ~~wherein R⁴ is optionally substituted with one or more groups selected from~~ C₁₋₆alkyl, halogenated C₁₋₆alkyl, -NO₂, -CF₃, C₁₋₆alkoxy, chloro, fluoro, bromo, and iodo;

R² and R³ are ethyl; R², R³, and

R⁴ is methyl ~~are, independently, C₁₋₃alkyl or halogenated C₁₋₃alkyl; and~~

R⁵ is hydrogen.

4. (currently amended) A compound according to claim 1, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof

wherein R¹ is ~~selected from~~ phenyl, pyridyl, thienyl, furyl, imidazolyl, pyrrolyl, ~~and or~~ thiazolyl;

R² and R³ are ethyl;

R⁴ is C₁₋₃alkyl; and

R⁵ is hydrogen.

5. (currently amended) A compound according to claim 1, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof, wherein the compound is selected from:

N,N-diethyl-4-[[3-[(methylsulfonyl)amino]phenyl][1-(thien-2-ylmethyl)piperidin-4-ylidene]methyl]benzamide;

N,N-diethyl-4-[[1-(2-furanylmethyl)-4-piperidinylidene][3-[(methylsulfonyl)amino]phenyl]methyl]benzamide;

N,N-diethyl-4-[[1-(phenylmethyl)-4-piperidinylidene][3-[(methylsulfonyl)amino]phenyl]methyl]-benzamide;

N,N-diethyl-4-[[3-[(methylsulfonyl)amino]phenyl][1-(3-pyridinylmethyl)-4-piperidinylidene]methyl]-benzamide; and

N,N-diethyl-4-[[3-[(methylsulfonyl)amino]phenyl][1-(3-thiazolyl-methyl)-4-piperidinylidene]methyl]-benzamide;

~~and pharmaceutically acceptable salts thereof.~~

6. (cancelled)

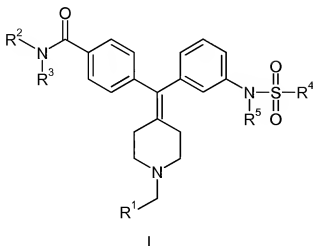
7. (currently amended) A method for the therapy of pain, anxiety or functional gastrointestinal disorders, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof.

8. (currently amended) A pharmaceutical composition comprising a compound according to claim 1, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof and a pharmaceutically acceptable carrier.

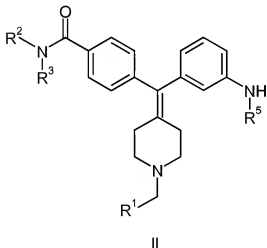
9. (currently amended) A method for the therapy of pain in a warm-blooded animal, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof.

10. (currently amended) A method for the therapy of functional gastrointestinal disorders in a warm-blooded animal, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof.

11. (currently amended) A process for preparing a compound of formula I, comprising:



reacting a compound of formula II with $X-S(=O)_2-R^4$ or $R^4S(=O)_2-O-S(=O)_2R^4$.



wherein

X is selected from Cl, Br and I;

R^1 is phenyl; pyridyl; thienyl; furyl; imidazolyl; pyrrolyl; or thiazolyl, wherein R^1 is optionally substituted with one or more groups selected from C_{1-6} alkyl, halogenated C_{1-6} alkyl, $-NO_2$, $-CF_3$, C_{1-6} alkoxy, chloro, fluoro, bromo, and iodo; selected from C_{6-10} aryl and C_{2-6} heteroaryl, wherein said C_{6-10} aryl and C_{2-6} heteroaryl are optionally substituted with one or more groups selected from $-R$, $-NO_2$, $-OR$, $-Cl$, $-Br$, $-I$, $-F$, $-CF_3$, $-C(=O)R$, $-C(=O)OH$, $-NH_2$, $-SH$, $-NHR$, $-NR_2$, $-SR$, $-SO_3H$, $-SO_2R$, $-S(=O)R$, $-CN$, $-OH$, $-C(=O)OR$, $-C(=O)NR_2$, $-NRC(=O)R$, and $-NRC(=O)-OR$, wherein R is, independently, a hydrogen or C_{1-6} alkyl; and

R^2 , R^3 , and R^4 and R^5 are, independently, C_{1-3} alkyl or halogenated C_{1-3} alkyl selected from hydrogen, C_{1-6} alkyl, and C_{3-6} cycloalkyl, wherein said C_{1-6} alkyl and C_{3-6} cycloalkyl are optionally substituted with one or more groups selected from $-R$, $-NO_2$, $-OR$, $-Cl$, $-Br$, $-I$, $-F$, $-CF_3$, $-C(=O)R$,

~~C(-O)CH₂-NH₂-SH, -NHR, -NR₂, -SR, -SO₃H, -SO₂R, -S(-O)R, -CN, -OH, C(=O)OR, C(=O)NR₂, -NRC(=O)R, and -NRC(=O)-OR, wherein R is, independently, a hydrogen or C₁₋₆alkyl; and~~
R⁵ is hydrogen.

12. (currently amended) A method for the therapy of pain, anxiety or functional gastrointestinal disorders, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 2, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof.

13. (currently amended) A method for the therapy of pain, anxiety or functional gastrointestinal disorders, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 3, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof.

14. (currently amended) A method for the therapy of anxiety, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 1, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof.

15. (currently amended) A method for the therapy of anxiety, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 2, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof.

16. (currently amended) A method for the therapy of anxiety, comprising ~~the step of~~ administering to said animal in need of such therapy a therapeutically effective amount of a compound according to claim 3, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof.

17. (currently amended) A pharmaceutical composition comprising a compound according to claim 2, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof and a pharmaceutically acceptable carrier.

18. (currently amended) A pharmaceutical composition comprising a compound according to claim 3, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof and a pharmaceutically acceptable carrier.

19. (currently amended) A pharmaceutical composition comprising a compound according to claim 4, or pharmaceutically acceptable salts thereof, or diastereomers, enantiomers, or mixtures thereof and a pharmaceutically acceptable carrier.